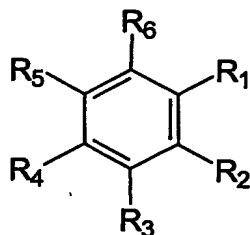


## CLAIMS

1. A method for modifying an amino-terminated surface of a solid support with carboxy groups comprising the steps of:

- 5       a) providing an amino-terminated surface; and  
      b) contacting the surface with a compound of the general formula (I):



(I)

10       wherein at least three of R<sub>1</sub> to R<sub>6</sub> are, independent from each other, selected from

15       - (CH<sub>2</sub>)<sub>n</sub> - (C=O) - X - Y - Z, and the remaining R groups are H;  
      or R<sub>1</sub> and R<sub>2</sub> form a ring, preferably an anhydride;

      X is a group selected from C<sub>1</sub>-C<sub>10</sub> alkyl, C<sub>1</sub>-C<sub>10</sub> alkenyl, a C<sub>3</sub>-C<sub>8</sub> cycloalkyl, aryl, heteroaryl, or a polyethylene glycol chain of the general form (CH<sub>2</sub>-CH<sub>2</sub>-O)<sub>m</sub>, wherein m is an integer from 1 to 450, or X is a bond;

20       Y is a carbonyl group, or a bond;

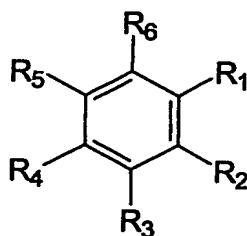
      Z is OH or an electron withdrawing group; and

      n is an integer from 0 to 10,

      under conditions allowing the formation of an amide bond between a carboxy group of the compound of the general  
25       formula (I) and the amino group of the solid surface.

2. A method for modifying an amino-terminated surface of a solid support with carboxy groups comprising the steps of:

- a) providing an amino-terminated surface; and
- b) contacting the surface with a compound of the general formula (I):



(I)

wherein:

(i) at least three of  $R_1$  to  $R_6$  are, independent from each other, selected from  $-(CH_2)_n-(C=O)-X-Y-Z$  and the remaining R groups are H; or

(ii)  $R_1$  and  $R_6$  are together of formula  $-(C=O)-Z'-(C=O)-$  so as to form a ring, at least one of  $R_2$  to  $R_5$  are, independent from each other, selected from

$-(CH_2)_n-(C=O)-X-Y-Z$  and the remaining R groups are H;

X is a group selected from  $C_1$ - $C_{10}$  alkyl,  $C_1$ - $C_{10}$  alkenyl, a  $C_3$ - $C_8$  cycloalkyl, aryl, heteroaryl, or a polyethylene glycol chain of the general form  $(CH_2-CH_2-O)_m$ , wherein m is an integer from 1 to 450, or X is a bond;

Y is a carbonyl group, or a bond;

Z is OH or an electron withdrawing group;

Z' is O or S; and

n is an integer from 0 to 10,

under conditions allowing the formation of an amide bond  
between a carboxy group of the compound of the general  
5 formula (I) and the amino group of the solid surface.

3. The method of claim 1 or claim 2, wherein the solid  
support is glass, a polymer, a metal, a semiconductor or an  
insulator.

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4. The claim of any one preceding claim wherein the solid  
support is glass, a polymer, a metal, a semiconductor or an  
insulator, particularly preferred the surface is an amine-  
terminated siloxane surface.

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5. The method of any one of claims 1 to 4, wherein  $n=0$  for  
the at least three of  $R_1$  to  $R_6$ , and the remaining R groups  
are H.

20

6. The method of any one of claims 1 to 4, wherein  $n=1$  for  
the at least three of  $R_1$  to  $R_6$ , and the remaining groups are  
H.

25

7. The method of any one of claims 1 to 4, wherein n is an  
integer from 2 to 5 for each of the at least three of  $R_1$  to  
 $R_6$ , and the remaining groups are H.

8. The method as described in any one preceding claim wherein three of  $R_1$  to  $R_6$  are, independent from each other, selected from  $-(CH_2)_n-(C=O)-X-Y-Z$  and the remaining R groups are H.

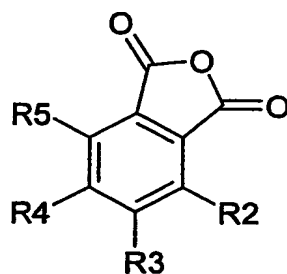
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9. The method as claimed in claim 8, wherein two of  $R_1$  to  $R_6$  are of formula  $-(CH_2)_n-(C=O)-X-Y-Z$  and have  $Z=\text{electron withdrawing group}$  and the remaining R group is of formula  $-(CH_2)_n-(C=O)-X-Y-Z$  and has  $Z=OH$ ; or two of  $R_1$  to  $R_6$  are of  
10 formula  $-(CH_2)_n-(C=O)-X-Y-Z$  and have  $Z=OH$  and the remaining R group is of formula  $-(CH_2)_n-(C=O)-X-Y-Z$  and has  $Z=\text{electron withdrawing group}$ .

10. The method of any one preceding claim wherein the at  
15 least three of  $R_1$  to  $R_6$  are  $R_1$ ,  $R_3$  and  $R_5$ .

11. The method as claimed in claim 2, wherein  $Z'$  is O such that said ring is an anhydride.

20 12. The method as claimed in claim 1 or claim 11, wherein the compound of general formula (I) is of the general formula (II):



(II)

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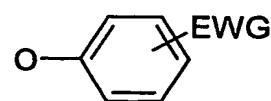
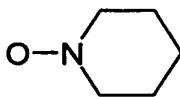
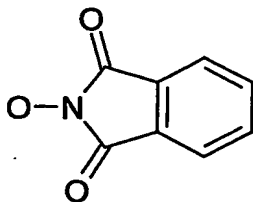
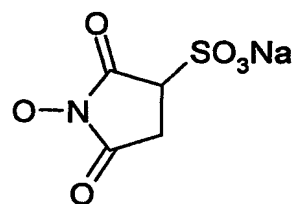
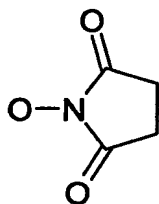
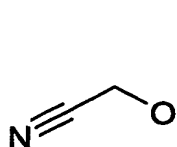
13. The method as claimed in any one preceding claim wherein Z is a leaving group, selected so as to make an activated derivative of carboxylic acid.

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14. The method as claimed in any one preceding claim, wherein Z is a halogen atom selected from F, Cl and Br.

15. The method as claimed in any one of claims 1 to 13,  
10 wherein Z is selected from phenol substituted by at least one strong electron withdrawing group, cyanomethyl, hydroxysuccinimide or its sodium sulfonate derivative NHS or sulfo-NHS, hydroxyphthalimide, and hydroxypiperidine.

15 16. The method as claimed in claim 15, wherein Z is selected from:



wherein EWG is an electron withdrawing group.

17. The method as claimed in any of claims 1 to 10, wherein the compound is trimesic acid or a mono-, di-, or tri-succinimidyl ester thereof, or the compound is benzene-1,3,5-triacetic acid or a mono-, di-, or tri-succinimidyl ester thereof.

18. The method as claimed in claim 17, wherein the compound is trimesic acid or a mono- or di-succinimidyl ester thereof, or the compound is benzene-1,3,5-triacetic acid or a mono- or di-succinimidyl ester thereof.

19. The method as claimed in claim 17, wherein the compound is a tri-succinimidyl ester of trimesic acid, or the compound is a tri-succinimidyl ester of benzene-1,3,5-triacetic acid.

20. The method as claimed in any one of claims 17 to 19, wherein said succinimidyl ester is a substituted succinimidyl ester.

21. The method of claim 20 wherein said substituted succinimidyl ester is a sulfonate derivative of succinimide.

22. The method as claimed in any one preceding claim wherein a coupling reagent is present.

23. The method as claimed in claim 22, wherein the coupling reagent comprises an uronium- or phosphonium-based coupling reagent.

24. The method as claimed in claim 23, wherein the coupling reagent comprises benzotriazol-1-yloxytris (dimethylamino)phosphonium hexafluorophosphate (BOP).

5 25. The method as claimed in claim 22, wherein the coupling reagent comprises a carbodiimide.

26. The method as claimed in claim 25, wherein the carbodiimide is dicyclohexylcarbodiimide,  
10 diisopropylcarbodiimide or 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride.

27. The method as claimed in claim 26, wherein the carbodiimide is 1-(3-dimethylaminopropyl)-3-  
15 ethylcarbodiimide hydrochloride.

28. The method as claimed in any one preceding claim wherein in step b) an excess of a tertiary base is added.

20 29. The method as claimed in claim 28, wherein the tertiary base is diisopropylethylamine, triethylamine, N-ethylmorpholine or N-methylmorpholine.

30. The method as claimed in any one preceding claim  
25 wherein in step b) additionally hydroxycinnamic acid is added.

31. The method as claimed in any one preceding claim wherein in step b) the amount of the compound is limiting,  
30 preferably such that not all amino groups of the solid support are carboxylated.

32. A solid surface obtainable by the method according to any of claims 1 to 31.

33. The solid surface of claim 32 which is a carboxy-terminated solid surface.

34. A method for conjugating an amino-group containing substrate to an amino-terminated surface of a solid support comprising:

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i) performing the steps as defined in any of claims 1 to 31 to obtain a carboxy-terminated surface of a solid support; and

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ii) contacting the amino-group containing substrate with the carboxy-terminated surface of the solid support of step i) under conditions allowing the formation of an amide bond between the carboxy group of the surface of the solid support and the amino group of the amino-group-containing substrate.

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35. The method as claimed in claim 34, wherein in step ii) a coupling reagent as defined in any of claims 22 to 27 is present.

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36. The method as claimed in claim 34 or claim 35, wherein in step ii) a tertiary base as defined in either claim 28 or claim 29 is present.

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37. The method as claimed in any one of claims 34 to 36, wherein the amino group-containing substrate is derived from

nucleotides, amino acids, sugars, oligomers or polymers thereof.

38. A compound of the general formula (I) as defined in any  
5 one of claims 1 to 21.

39. A method for preparing the compound according to any of  
claims 1 to 21 comprising preparing the compound in a manner  
known *per se*.